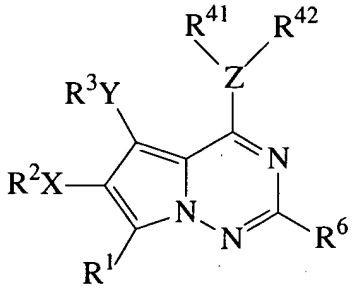
AMENDMENTS TO THE CLAIMS

Please amend the claims as follows:

Claim 1 (canceled). A compound of formula (I)



(I)

wherein

Z is selected from the group consisting of O, S, N, OH, and Cl, with the provisos that when Z is O or S, R^{41} is absent, and when Z is OH or Cl, both R^{41} and R^{42} are absent, and when Z is N, then R^{41} is H;

X and Y are independently selected from the group consisting of O, OCO, S, SO, SO_2, CO, CO_2, NR^{10}, $NR^{11}CO$, $NR^{12}CONR^{13}$, $NR^{14}CO_2$, $NR^{15}SO_2$, $NR^{16}SO_2NR^{17}$, SO_2NR^{18}, $CONR^{19}$, halogen, nitro and cyano, or X or Y are absent;

R^1 is hydrogen, CH_3, OH, OCH_3, SH, SCH_3, $OCOR^{21}$, SOR^{22}, SO_2R^{23}, $SO_2NR^{24}R^{25}$, CO_2R^{26}, $CONR^{27}R^{28}$, NH_2, $NR^{29}SO_2NR^{30}R^{31}$, $NR^{32}SO_2R^{33}$, $NR^{34}COR^{35}$, $NR^{36}CO_2R^{37}$, $NR^{38}CONR^{39}R^{40}$, halogen, nitro, or cyano;

R^2 and R^3 are independently hydrogen, alkyl, substituted alkyl, alkenyl, substituted alkenyl, alkynyl, substituted alkynyl, aryl, substituted aryl, heterocyclo, substituted heterocyclo, aralkyl, substituted aralkyl, heteroaryl, substituted heteroaryl, heterocycloalkyl or substituted heterocycloalkyl; with the proviso that when X is halo, nitro or cyano, R^2 is absent, and, when Y is halo, nitro or cyano, R^3 is absent;

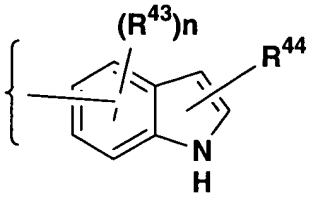
R^6 is H, alkyl, substituted alkyl, aryl, substituted aryl, heterocyclo, substituted heterocyclo, NR^7R^8, OR^9 or halogen;

R^7, R^8, R^9, R^{10}, R^{11}, R^{12}, R^{13}, R^{14}, R^{15}, R^{16}, R^{17}, R^{18}, R^{19}, R^{21}, R^{24}, R^{25}, R^{26}, R^{27}, R^{28}, R^{29}, R^{30}, R^{31}, R^{32}, R^{34}, R^{35}, R^{36}, R^{38}, R^{39} and R^{40} are independently selected from the

group consisting of hydrogen, alkyl, substituted alkyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocyclo, or substituted heterocyclo;

R^{22}, R^{23}, R^{33} and R^{37} are independently selected from the group consisting of alkyl, substituted alkyl, aryl, substituted aryl, heteroaryl, substituted heteroaryl, heterocyclo, or substituted heterocyclo;

R^{42} is



$(R^{43})_n$ wherein n equals 0, 1 or 2 and each R^{43} is independently selected from the group consisting of hydrogen, fluorine, chlorine and methyl; and

R^{44} is methyl, or hydrogen,

with the further provisos that:

a. R^2 may not be hydrogen if X is SO, SO_2, $NR^{13}CO_2$, or $NR^{14}SO_2$; and

b. R^3 may not be hydrogen if Y is SO, SO_2, $NR^{13}CO_2$, or $NR^{14}SO_2$;

or an enantiomer, diastereomer, or pharmaceutically acceptable salt, prodrug, or solvate thereof,

Claim 2 (canceled). A compound according to claim 1 wherein R^1 is hydrogen or methyl; R^6 is hydrogen; R^3 is lower alkyl; and Z is oxygen or nitrogen.

Claim 3 (canceled). A compound according to claim 1 wherein R^1 is hydrogen; R^3 is lower alkyl; Y is absent; X is oxygen or nitrogen; R^{43} is fluoro or hydrogen; and R^{44} is hydrogen or methyl.

Claim 4 (canceled). A compound according to claim 1 wherein X is oxygen; R^2 is a substituted alkyl and R^{43} is fluoro.

Claim 5 (Currently amended). A compound or a pharmaceutically acceptable salt thereof selected from the group consisting of

4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-ol,

1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methy-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-4-(aminosulfonyl)aminobutan-2-ol,

N-{3-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methy-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-2-Hydroxy-propyl}-methanesulfonamide,

(2*S*)-3-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methy-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propane-1,2-diol,

(2*R*)-3-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propane-1,2-diol,

(2*R*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propan-2-ol,

(2*S*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propan-2-ol,

(2*R*)1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-3-methoxy-propan-2-ol,

(2*S*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-3-methoxy-propan-2-ol,

2-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-ethanol,

N-{2-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methyl-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-ethyl}-methanesulfonamide,

(2*R*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methy-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-4-methanesulfonyl-butan-2-ol,

(2*S*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methy-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-4-methanesulfonyl-butan-2-ol,

5-Methyl-4-(2-methyl-*1H*-indol-5-yloxy)-6-(3-piperidin-1-ylpropoxy)-pyrrolo[2,1-f][1,2,4]triazine,

4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methyl-6-(2-piperidin-4-yl-ethoxy)-pyrrolo[2,1-f][1,2,4]triazine,

4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methyl-6-(3-pyridin-4-yl-propoxy)-pyrrolo[2,1-f][1,2,4]triazine,

{1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methyl-pyrrolo[2,1-f][1,2,4]triazin-6-yloxymethyl]-3-methanesulfonyl-propyl}-dimethyl-amine,

2-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-1-methylethylamine,

{2-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-1-methylethyl}-methylamine,

4-(4-Fluoro-2-methyl-1*H*-indol-5-yloxy)-5-methyl-6-(morpholin-2-ylmethoxy)-pyrrolo[2,1-f][1,2,4]triazine,

[(1*R*),2*S*]-2-Dimethylaminopropionic acid-[2-[4-(4-fluoro-2-methyl-1*H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f]-[1,2,4]triazin-6-yloxy]]-1-methylethyl ester,

[(1*R*), 2*S*]-2-Amino-4-methylpentanoic acid [2-[4-(4-fluoro-2-methyl-1H-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]]-1-methylethyl ester,

[(1*R*), 2*S*]-2-Aminopropionic acid 2-[4-(4-fluoro-2-methyl-1H-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-1-methylethyl ester,

4-(4-Fluoro-2-methyl-1H-indol-5-yloxy)-6-(3-methanesulfonyl-propoxy)-5-methyl-pyrrolo[2,1-f][1,2,4]triazine, and

N-{3-[4-(4-Fluoro-2-methyl-1H-indol-5-yloxy)-5-methyl-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propyl}-methanesulfonamide.

Claim 6 (Currently amended). A compound <u>or a pharmaceutically acceptable salt thereof</u> selected from the group consisting of

4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-ol,

(2*S*)-3-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methy-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propane-1,2-diol,

(2*R*)-3-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propane-1,2-diol,

(2*R*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propan-2-ol,

(2*S*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propan-2-ol,

(2*R*)1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-3-methoxy-propan-2-ol,

(2*S*)-1-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-3-methoxy-propan-2-ol,

5-Methyl-4-(2-methyl-*1H*-indol-5-yloxy)-6-(3-piperidin-1-ylpropoxy)-pyrrolo[2,1-f][1,2,4]triazine,

4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methyl-6-(2-piperidin-4-yl-ethoxy)-pyrrolo[2,1-f][1,2,4]triazine,

2-[4-(4-Fluoro-2-methyl-*1H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-1-methylethylamine,

[(1*R*),2*S*]-2-Dimethylaminopropionic acid-[2-[4-(4-fluoro-2-methyl-1*H*-indol-5-yloxy)-5-methylpyrrolo[2,1-f]-[1,2,4]triazin-6-yloxy]]-1-methylethyl ester,

[(1*R*), 2*S*]-2-Amino-4-methylpentanoic acid [2-[4-(4-fluoro-2-methyl-1H-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]]-1-methylethyl ester,

[(1*R*), 2*S*]-2-Aminopropionic acid 2-[4-(4-fluoro-2-methyl-1H-indol-5-yloxy)-5-methylpyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-1-methylethyl ester,

4-(4-Fluoro-2-methyl-1H-indol-5-yloxy)-6-(3-methanesulfonyl-propoxy)-5-methyl-pyrrolo[2,1-f][1,2,4]triazine, and

N-{3-[4-(4-Fluoro-2-methyl-1H-indol-5-yloxy)-5-methyl-pyrrolo[2,1-f][1,2,4]triazin-6-yloxy]-propyl}-methanesulfonamide.

Claim 7 (canceled). A pharmaceutical composition comprising at least one of the compounds of Claim 1 and a pharmaceutically acceptable carrier therefor.

Claim 8 (Original). A pharmaceutical composition comprising at least one of the compounds of Claim 5 and a pharmaceutically acceptable carrier therefor.

Claim 9 (Original). A pharmaceutical composition comprising at least one of the compounds of Claim 6 and a pharmaceutically acceptable carrier therefor.

Claim 10 (Canceled). A pharmaceutical composition comprising at least one or more compounds of Claim 1 in combination with a pharmaceutically acceptable carrier and at least one additional anti-cancer or cytotoxic agent.

Claim 11 (Original). A pharmaceutical composition comprising at least one or more compounds of Claim 5 in combination with a pharmaceutically acceptable carrier and at least one additional anti-cancer or cytotoxic agent.

Claim 12 (Original). A pharmaceutical composition comprising at least one or more compounds of Claim 6 in combination with a pharmaceutically acceptable carrier and at least one additional anti-cancer or cytotoxic agent.

Claim 13 (Original). The pharmaceutical composition of Claim 8, wherein said anti-cancer or cytotoxic agent is selected from the group consisting of: linomide, inhibitors of integrin $\alpha v\beta 3$ function, angiostatin, razoxane, tamoxifen, toremifene, raloxifene, droloxifene, iodoxifene, megestrol acetate, anastrozole, letrozole, borazole, exemestane, flutamide, nilutamide, bicalutamide, cyproterone acetate, gosereline acetate, leuprolide, finasteride, herceptin, metalloproteinase inhibitors, inhibitors of urokinase plasminogen activator receptor function, growth factor antibodies, growth factor receptor antibodies, bevacizumab, cetuximab, tyrosine kinase inhibitors, serine/threonine kinase inhibitors, methotrexate, 5-fluorouracil, purine, adenosine analogues, cytosine arabinoside, doxorubicin, daunomycin, epirubicin, idarubicin, mitomycin-C, dactinomycin, mithramycin, cisplatin, carboplatin, nitrogen mustard, melphalan, chlorambucil, busulphan, cyclophosphamide, ifosfamide, nitrosoureas, thiotepa, vincristine, paclitaxel, docetaxel, epothilone analogs, discodermolide analogs, eleutherobin analogs, etoposide, teniposide, amsacrine, topotecan, irinotecan, flavopyridols, proteasome inhibitors including bortezomib and biological response modifiers.

Claim 14 (canceled). A method for producing an antiangiogenic effect which comprises administering to a mammalian species in need thereof, an effective antiangiogenic producing amount of at least one of the compounds of Claim 1.

Claim 15 (canceled). A method for producing a vascular permeability reducing effect which comprises administering to a mammalian species in need thereof an effective vascular permeability reducing amount of at least one of the compounds of Claim 1.

Claim 16 (canceled). A method of inhibiting protein kinase activity of growth factor receptors which comprises administering to a mammalian species in need thereof, an effective protein kinase inhibiting amount of at least one of the compounds of Claim 1.

Claim 17 (canceled). A method of inhibiting tyrosine kinase activity of growth factor receptors which comprises administering to a mammalian species in need thereof, an effective tyrosine kinase inhibiting amount of at least one of the compounds of Claim 1.

Claim 18 (canceled). A method for treating proliferative diseases, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim7.

Claim 19 (canceled). A method for treating cancer, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 7.

Claim 20 (canceled). A method for treating inflammation, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 7.

Claim 21 (canceled). A method for treating autoimmune diseases, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 7.

Claim 22 (canceled). A method for treating proliferative diseases, comprising administering to mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 7.

Claim 23 (canceled). A method for treating cancer, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 7.

Claim 24 (canceled). A method for treating inflammation, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 7.

Claim 25 (canceled). A method for treating autoimmune diseases, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 7.

Claim 26 (canceled). A method for treating diseases associated with signal transduction pathways operating through growth factor receptors, which comprises administering to a mammalian species in need thereof a therapeutically effective amount of at least one of the compounds of Claim 1.

Please add the following new claims:

Claim 27 (new). A method for treating proliferative diseases, comprising administering to mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 8.

Claim 28 (new). A method for treating cancer, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 8.

Claim 29 (new). A method for treating inflammation, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 8.

Claim 30 (new). A method for treating autoimmune diseases, comprising administering to a mammalian species in need thereof, a therapeutically effective amount of the composition of Claim 8.

Claim 31 (new). A method for treating diseases associated with signal transduction pathways operating through growth factor receptors, which comprises administering to a mammalian species in need thereof a therapeutically effective amount of at least one of the compounds of Claim 8.